What is claimed is:

1. A compound selected from the group of compounds represented by Formula (I):

(I)

5 wherein:

R¹ and R⁴ are, independently of each other, hydrogen or alkyl;

R² is: (i) cycloalkyl, cycloalkylalkyl, aryl, aralkyl, aralkenyl, heteroaryl, heteroaralkyl, heteroaralkyl, heteroaralkyl, or

(ii) -(alkylene)-B-X where B is -O-, -NR 8 -, -S(O)_n- (where n is 0, 1 or 2), -C=O, -CONR 8 -, -NR 8 CO₂-, NR 8 SO₂- or -C(=NR 8)NR 8 SO₂-(where R 8 is H or alkyl), and X is cycloalkyl, cycloalkyl, aryl, aralkyl heteroaryl or heteroaralkyl; or

(iii) -(alkylene)-B-X where B is -NR⁸CO- (where R⁸ is H or alkyl), and X is cycloalkyl, cycloalkylalkyl, aryl, aralkyl heteroaryl or heteroaralkyl; or

(iv) R² and R³ form an alkylene or heteroalkylene chain;

R³ is hydrogen or alkyl;

R⁶ is hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, aryl, aralkyl, heteroaryl or heteroaralkyl;

R⁵ is:

(i) hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, aryl, aralkyl, aralkenyl, heteroaryl, 20 heteroaralkyl, heteroaralkenyl, heterocycloalkyl, heteroalkyl, or -(alkylene)-C(O)-X¹ where X¹ is alkyl, hydroxy, alkoxy, aryl, aralkyl, aryloxy, aralkyloxy, heteroaryl, heteroaryloxy, heteroaralkyloxy or NR'R" (where R' and R" are independently H or alkyl, or R'and R" form an alkylene chain); or

(ii) R⁵ and R⁴ form an alkylene chain; or

(iii) R⁵ and R⁶ form an alkylene chain;

n is 0 or 1;

A is $-C(=O)-CH(R^9)-(CH_2)_m-N(R^{10})$ - wherein:

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 R^9 is hydrogen, alkyl, cycloalkyl, cycloalkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heteroaryl, or -(alkylene)-C(O)- X^1 where X^1 is alkyl, hydroxy, alkoxy, aryl, aralkyl, aryloxy, aralkyloxy, heteroaryl, heteroaryloxy, heteroaralkyloxy or NR'R"

5 (where R' and R" are independently H or alkyl, or R' and R" form an alkylene chain); and R¹⁰ is hydrogen, alkyl, aralkyl or heteroaralkyl;

Z is Y-B wherein:

Y is alkylene or a bond; and B is -CO-, -C(O)O-, -CONR⁸-, -SO₂-, or -SO₂NR⁸- (where R⁸ is hydrogen or alkyl), alkylene (optionally substituted by hydroxy, alkoxy, amino, monoalkylamino or dialkylamino) or a bond;

 R^7 is cycloalkyl, cycloalkylalkyl, aryl, aralkyl, heteroaryl or heteroaralkyl; provided that when n=0 and Z is SO_2 , then R^2 does not contain an imidazole group; and their pharmaceutically acceptable salts, prodrugs, individual isomers, and mixtures of isomers.

- 2. The compound of Claim 1 wherein: n is 0.
- 3. The compound of Claim 2 wherein R^3 and R^6 are hydrogen.
- 4. The compound of Claim 3, wherein:

 R² is aralkyl or heteroaralkyl.
- 5. The compound of Claim 4 wherein: Z is -C(O)O- or S(O)₂-.
- The compound of Claim 5 wherein:R² is optionally substituted benzyl or heteroaralkylmethyl.

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- The compound of Claim 6 wherein, R² is 4-t-butoxybenzyl, 3-chlorobenzyl, 3-indolyl 7. methyl, 2-thienylmethyl, 4-imidazolylmethyl or 4-thiazolylmethyl.
- The compound of Claim 7 wherein R² is 4-thiazolylmethyl. 8.

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The compound of Claim 8 wherein: 9. R⁷ is aryl, aralkyl, heteroaryl or heteroaralkyl.

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The compound of Claim 8 wherein:

Z is -C(O)O- and \mathbb{R}^{7} is optionally substituted benzyl.

The compound of Claim 9 wherein: 11.

Z is -SO₂- and R⁷ is aryl or heteroaryl.

The compound of Claim 10, wherein: 12.

R¹ and R⁴ are hydrogen and R⁵ is alkyl.

- The compound of Claim 12 wherein R⁵ is (S,S)-1-methylpropyl. 13.
- 20 14. The compound of Claim 11, wherein: R¹ and R⁴ are hydrogen and R⁵ is alkyl.
 - The compound of Claim 14 wherein R⁵ is (S,S)-1-methylpropyl. 15.

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The compound of Claim 3 wherein:

R² is (alkylene)-B-X where B is -O-, -NR⁸-, -S(O)_n- (where n is 0, 1 or 2), -C=O, -CONR⁸-, -NR⁸CO₂-, -NR⁸SO₂- on -C = NR⁸)NSO₂-(where R⁸ is H or alkyl), and X is cycloalkyl, cycloalkylalkyl, aryl, aralkyl heteroaryl or heteroaralkyl.



The compound of Claim 16, wherein: **17.**

Z is -C(O)O- or $-S(O)_2$ -.

- The compound of Claim 17, wherein R² is CH₂-B-X and 18. B is -NHCO₂- and X is benzyl.
- 19. The compound of Claim 18 wherein: R⁷ is aryl or aralkyl.

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The compound of Claim 19, wherein: 20.

R¹ and R⁴ are hydrogen and R⁵ is alkyl.

The compound of Claim 20 wherein R⁵ is (S,S)-1-methylpropyl. 21.

The compound of Claim 1 wherein: 22. 15 n is 1.

- The compound of Claim 22 wherein m is 0 and R³ and R⁶ are hydrogen. 23.
- The compound of Claim 23, wherein: 20 24. R² is aralkyl or heteroaralkyl.

The compound of Claim 24, wherein:

Z is $-C(O)O - or -S(O)_2$ -.

- The compound of Claim 25, wherein: 26.
 - R² is optionally substituted benzyl or heteroarylmethyl.
 - The compound of Claim 26 wherein R² is 4-t-butoxybenzyl, 3-chlorobenzyl, 3-indolyl 27. methyl, 2-thienylmethyl, 4-imidazolylmethyl or 4-thiazolylmethyl.

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- 28. The compound of Claim 27 wherein R² is 4-thiazolylmethyl.
- 29. The compound of Claim 28 wherein:R⁷ is aryl, aralkyl, heteroaryl or heteroaralkyl.

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- 30. The compound of Claim 29 wherein: Z is -C(O)O- and R⁷ is benzyl.
- **31.** The compound of Claim 29 wherein:

Z is $-SO_2$ - and R^7 is aryl.

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32. The compound of Claim 30, wherein: $R^{1} \text{ and } R^{4} \text{ are hydrogen and } R^{5} \text{ is alkyl.}$

- 15 33. The compound of Claim 32 wherein R⁵ is (S,S)-1-methylpropyl.
 - 34. The compound of Claim 31, wherein: R¹ and R⁴ are hydrogen and R⁵ is alkyl.
- 20 35. The compound of Claim 34 wherein R⁵ is (S,S)-1-methylpropyl.

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36. The compound of Claim 23, wherein:

R² is (alkylene)-B-X where B is -O-, -NR⁸-, -S-, -C=O, -CONR⁸-, -NR⁸CO₂-, -NSO₂- or -C(=NR⁸)NSO₂-(where R⁸ is Hor alkyl), and X is cycloalkyl, cycloalkylalkyl, aryl, aralkyl heteroaryl or heteroaralkyl)

37. The compound of Claim 36, wherein: Z is -C(O)O- or -S(O)₂-.



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- 38. The compound of Claim 37, wherein R² is CH₂-B-X and B is -NHCO₂- and X is benzyl.
- 39. The compound of Claim 38 wherein:R⁷ is aryl or aralkyl.
- 40. The compound of Claim 39, wherein:

 R¹ and R⁴ are hydrogen and R⁵ is alkyl.
- The compound of Claim 40 wherein R⁵ is (S,S)-1-methylpropyl.
 - 42. A pharmaceutical composition comprising the compound of Claim 1 and a pharmaceutically acceptable excipient.
 - 15 43. A method of treating disease comprising administering to a patient in need thereof a compound of Claim 1.
 - 44. The method of Claim 43, wherein the disease is a fibrotic disease.
 - 20 45. The method of Claim 44 wherein the disease is acute respiratory distress syndrome.
 - 46. A method of treating fibrosis comprising administering to a patient in need thereof an inhibitor of procollagen C-proteinase that is at least ten-fold more selective for procollagen C-proteinase over both collagenase-1, collagenase-2 and collagenase-3.

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- 47. A method for preparing the compounds of Claim 1 by:
- (i) treating a compound of Formula II wherein L is a leaving group and R^1 R^7 , A, n and Z are as defined in Claim 1 with hydroxylamine or a protected derivative thereof, and
 - (ii) deprotecting as necessary and isolating the compound of Claim 1.

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